

## CLAIMS

1. A fast-dissolving pharmaceutical composition comprising micronized (R)-2-(4-bromo-2-fluorobenzyl)-  
5 1,2,3,4-tetrahydropyrrolo[1,2-a]pyrazine-4-spiro-3'-pyrrolidine-1,2',3,5'-tetrone (hereinafter, referred to as "AS-3201").

2. The fast-dissolving pharmaceutical composition according to claim 1, wherein the mean particle size of the  
10 micronized AS-3201 is less than about 10  $\mu\text{m}$ .

3. The fast-dissolving pharmaceutical composition according to claim 1, wherein the mean particle size of the micronized AS-3201 is less than about 5  $\mu\text{m}$ .

4. The fast-dissolving pharmaceutical composition  
15 according to claim 1, wherein the mean particle size of the micronized AS-3201 is in the range of about 0.5  $\mu\text{m}$  - about 3  $\mu\text{m}$ .

5. The fast-dissolving pharmaceutical composition according to any one of claims 1-4, which comprises the  
20 micronized AS-3201 in a ratio of about 0.5 % by weight - 5 % by weight, a diluent in a ratio of about 51 % by weight - about 93.8 % by weight, a disintegrator in a ratio of about 5 % by weight - about 35 % by weight, a binder in a ratio of about 0.5 % by weight - about 5 % by weight, and a  
25 lubricant in a ratio of about 0.2 % by weight - about 4 %

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by weight, to the total weight of the pharmaceutical composition.

6. The fast-dissolving pharmaceutical composition according to claim 5, which comprises a diluent in a ratio of about 59 % by weight - about 88 % by weight, a disintegrator in a ratio of about 10 % by weight - about 30 % by weight, a binder in a ratio of about 1 % by weight - about 3 % by weight, and a lubricant in a ratio of about 0.5 % by weight - about 3 % by weight.

7. The fast-dissolving pharmaceutical composition according to any one of claims 1-4, which comprises the micronized AS-3201 in a ratio of more than 5 % by weight and less than about 25% by weight, a diluent in a ratio of about 16 % by weight - about 84.3 % by weight, a disintegrator in a ratio of about 10 % by weight - about 50 % by weight, a binder in a ratio of about 0.5 % by weight - about 5 % by weight, and a lubricant in a ratio of about 0.2 % by weight - about 4 % by weight, to the total weight of the pharmaceutical composition.

8. The fast-dissolving pharmaceutical composition according to claim 7, which comprises a diluent in a ratio of about 29 % by weight - about 73.5 % by weight, a disintegrator in a ratio of about 20 % by weight - about 40 % by weight, a binder in a ratio of about 1 % by weight - about 3 % by weight, and a lubricant in a ratio of about

0.5 % by weight - about 3 % by weight.

9. The fast-dissolving pharmaceutical composition according to any one of claims 1-8, which has a dissolution percentage of the active substance 50 % or more for 15 minutes after the start of the dissolution test.

10. The fast-dissolving pharmaceutical composition according to claim 9, which has a dissolution percentage of the active substance of 80 % or more for 15 minutes after the start of the dissolution test.

11. The fast-dissolving pharmaceutical composition according to any one of claims 1-10, which contains as a stabilizer at least one acidic substance having an acidity more potent than that of AS-3201.

12. The fast-dissolving pharmaceutical composition according to claim 11, wherein the acidic substance is a member selected from the group consisting of citric acid, tartaric acid, maleic acid and phosphoric acid.

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